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(54) **METAL PIGMENT COMPOSITION**
METALLPIGMENTZUSAMMENSETZUNG
COMPOSITION DE PIGMENT METALLIQUE

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Description

[0001] The present invention relates to a process for preparing low- or non-dusting, substantially non-volatile highly light reflective metal pigment compositions by a ball milling process using water as the major milling fluid.

[0002] The preparation of metal flake pigments is well documented in the patent literature. For example, US Patent 3901688 describes a wet ball milling process in which metal powder or chopped foil is milled with an organic liquid such as mineral spirits and a small amount of a lubricant such as stearic or oleic acid. The metal flakes so produced are separated, for example by wet sieving to provide the desired particle size distribution and thereafter brought to a paste-like consistency of, typically, 55-80% weight metal content. In this conventional process the lubricant is required to avoid cold welding of the metal flakes under the action of the grinding media. The purpose of the organic liquid is to reduce the viscosity of the system, enabling efficient conversion of the starting powder to flakes.

[0003] Metal flakes may also be prepared in the complete absence of solvent by a dry ball milling process, such as that described in US Patent 4115107.

[0004] A further method of preparation of metal flake is by electrodeposition of a thin film of metal onto an inert support, followed by removal and fragmentation into flakes. The product is generally provided either as dry flake or as a dispersion of the metal flakes in solvent.

[0005] The aforementioned methods for preparing metal flake pigments suffer from a number of disadvantages. For example, dry milling processes are now little used, due to the explosive and contaminating properties of the dry flake products. Though the paste form, in which the metal flake particles are damped by solvent, is inherently safer and easier to handle, it is not without problems. The 20-45% by weight solvent portion of metal pigment pastes may be acceptable in certain coating applications such as automotive paints, but in others, especially printing inks, such solvents typically slow down the drying rate and may cause odour in the final printed film, due to retention of minute concentrations of these typically high boiling point solvents. This is especially disadvantageous in printed films on packaging intended for food contact. The presence of organic solvent in other application areas, such as plastics, can also be very undesirable. The solvent vaporises during processing, causing bubbles and surface blemishes in the pigmented plastic article.

[0006] Flake formation by the electrodeposition process can give very bright flakes of narrow particle size distribution, but it is a very costly process, unsuited to large scale production. This is because the equipment required to maintain the high vacuum environment in which the metal is vaporised is very expensive to construct and operate.

[0007] It is also found that the storage stability of the metal pigment pastes themselves is finite, due to the

tendency of the organic solvent component to evaporate, leading to aggregation of the metal flake particles. This is especially true for pastes in which a more highly volatile solvent has been employed, perhaps to ensure the compatibility of the metal pigment paste in a particular coating system. Once aggregated, redispersion is difficult. In derived coatings, hiding power, or opacity, and also gloss, may be greatly reduced.

[0008] Increasingly too, environmental concern about organic solvent is leading to legislation which encourages a lower use of volatile solvents in industrial processes. The costs associated with solvent purchase, storage and recovery are increasing. Containment equipment, required to meet increasingly severe legislative constraints on the emission of solvent to the atmosphere is expensive. As a result replacement of organic solvents by water is a strategic goal in many industries.

[0009] Recent changes in health and safety classifications require a number of high boiling point petroleum derived hydrocarbon solvents traditionally used in metal pigment manufacture to be designated carcinogenic. This increases the pressure for their elimination from metal flake pigment manufacturing processes.

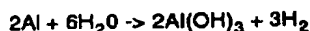
[0010] The need for a stable, safe and easily handled product metal pigment form, free of the disadvantages of both dry flakes and solvent containing pastes has been met by the products of European Patent 0134678. There is described therein a process for the preparation of a solid low- or non-dusting, metal pigment composition which comprises forming a coherent paste of an organic binder medium, an organic liquid vehicle and metal pigment, in powder or flake form, the paste being formed by mixing a first component comprising organic binder medium and a second component comprising metal pigment, with either or both of the first and second components comprising organic liquid vehicle and the paste containing from 3-45% of the organic binder medium based on the weight of the metal pigment, and either sub-dividing the coherent paste into particles and removing substantially all organic liquid vehicle from the particles, or removing substantially all the organic liquid vehicle from the coherent paste and subdividing the resulting mass into particles, at least 98% by weight of the resulting particles being retained on a sieve having a 150 μ m aperture and each containing a plurality of metal pigment particles dispersed in a matrix of organic binder medium. The so-called "granule" products of this process can be prepared using organic binder media compatible with the end application. Thus, for example, synthetic aldehyde and ketone resins could be employed for ink applications and polyolefin derivatives for the mass pigmentation of thermoplastics.

[0011] Though the products of European Patent 0134678 are substantially free of organic solvent, the process itself still involves processing of solvent. It is thus still subject to the aforementioned legislative pressures. There is therefore a clear need for a process for preparing an easily handled, dust free, metal pigment

product, which does not employ organic solvents.

[0012] It is possible to prepare a dust free, metal pigment product without the use of organic solvents if the metal pigment is prepared by dry milling and thereafter incorporated into a suitable organic carrier material by mixing at a temperature above the melting point of the organic carrier. Once a homogeneous mixture has been attained, the mass is cooled, often with extrusion into a more easily handled form, such as granules or pellets. Such processes were once widely employed for the preparation of plastics masterbatch. The need to use dry metal pigments, with their attendant disadvantages, and the difficulty of wetting such pigments satisfactorily into the carrier resin, have contributed to the decline in this type of process. Although the problem of metal flake dusting may be reduced if the metal powder precursor is milled in the presence of the carrier resin, methods traditionally used to separate wanted from unwanted particle size fraction, such as dilution with solvent and screening, are then no longer operable.

[0013] A more satisfactory processing liquid is water. It is readily available, inexpensive and non-flammable. However, a difficulty arises in the case of the more reactive metal pigments, especially aluminium, which is the most widely used of such pigments. Aluminium metal may react with water to form aluminium oxide and hydrogen gas, according to the equation:-



[0014] Generation of hydrogen is dangerous, because it is highly explosive. Sealed containers of aqueous metal pigment pastes may become pressurised and explode. In addition, the pigmentary properties of the metal flake are destroyed.

[0015] Many attempts have been made to stabilise aluminium against this reaction. Methods employed fall into several categories, for example resin encapsulation or chemical treatment. As an example of the former, there may be mentioned United States Patent 4213886. Prominent types of chemical treatment are silica coating (see, for example, United States Patent 2885386), treatment with chromium derivatives (German Patent 3636183), phosphate treatment, described in European Patent 0319971 and vanadium treatment (European Patent 0104075).

[0016] Milling aluminium powder in a mixture of water and an organic solvent has been described in United States Patent 3565655. Morpholine is used as a corrosion inhibitor with a fatty acid as the milling lubricant. However, the presence of at least 2 weight percent of a hydrocarbon or a halogenated hydrocarbon is a necessary part of the invention.

[0017] United States Patent 4693754 describes the milling of aluminium powder in a mixture of hydrocarbons, water, and a compound of chromium or vanadium which acts as a corrosion inhibitor. However again the

presence of some organic solvent in the milling process is a component part of the invention.

[0018] In WO-A-94/28074 there is described for the avoidance of odour the milling of aluminium powder, optionally with solvent or water, without a fatty acid lubricant, but in the presence of a polymer resin that is a paint or ink binder. This Patent Application does not include the use of a corrosion inhibitor in the milling process. In the absence of a corrosion inhibitor, water milled aluminium pigments are prone to severe gassing and result in products having a dull gray colour, in contrast to the bright metallic effects desired for decorative metallic finishes.

[0019] There is, therefore, still a need for a satisfactory process which can be used for milling aluminium powder and the like without organic solvent.

[0020] According to the present invention, there is provided a process for preparing a low- or non-dusting, substantially non-volatile (preferably highly light reflective) metal flake pigment composition, which comprises ball milling atomised metal powder in a milling fluid substantially comprised of water which has a maximum content of organic solvent of 10% by weight, in the presence of one or more additive(s) which together exhibit lubricating, corrosion inhibiting and surfactant properties, the lubricant being a non-suet lubricant.

[0021] Thereafter, unwanted oversize or undersize particles may be removed. In the next stage of the process the aqueous metal pigment slurry may be used unaltered, may be converted to a dry metal flake pigment powder by removal of the milling fluid, for example at elevated temperature, or is preferably concentrated to a paste-like consistency. The metal flake pigment can be incorporated in an organic binder medium by mixing to form a coherent paste of an organic binder medium, water and metal flake pigment, the paste being formed by mixing a first component comprising organic binder medium and a second component comprising metal pigment, with either or both of the first and second components comprising water and the paste containing from 1 to 150%; preferably 3-70%, and most preferably 30 to 60% of the organic binder medium based on the weight of the metal pigment, and either sub-dividing the coherent paste into particles and removing substantially all volatile liquids from the particles, or removing substantially all the volatile liquids from the coherent paste and sub-dividing the resulting mass into particles, at least 98% by weight of the resulting particles being retained on a sieve having 150 µm aperture and each containing a plurality of metal pigment particles dispersed in a matrix of organic binder medium, the organic binder medium being capable of binding the metal flake pigment particles by either precipitation from solution during volatile liquid removal, or melting or sintering at elevated temperature and fusion.

[0022] The volatile liquid will usually consist substantially of water. For example, the volatile liquid may be water together with small amounts of additives, for ex-

ample lower alcohols as wetting agents and resin solubilisers and dispersers.

[0023] The milling fluid used in the present invention consists substantially of water. The presence of up to 50% by volume (for example up to 30%, especially up to 25%, more especially up to 10%) of an organic solvent can usually be tolerated. A small amount (for example less than 2%) of an organic solvent is generally acceptable. Lower alcohols are especially useful as defoaming agents. Any such organic solvent is desirably water miscible. In one embodiment the milling fluid consists only of water and the system contains no organic liquid at all, the only organic compounds present being additives such as the lubricant(s), corrosion inhibitor(s) etc.

[0024] One of the advantages of the process of the present invention is the possibility of treating the milled flakes whilst in an aqueous system. For example, the milled flakes can be chromate treated or coated with silica or alumina prior to any granule formation. Such a process enables the pigments to be rendered suitable for different applications or to have different colour characteristics. Such aqueous treatments are known in the art, but tend to be difficult and expensive to carry out since the traditional milling fluid must be removed before the pigments can be treated in an aqueous medium. Since the milling process of the present invention is carried out in water, such treatments are simple to conduct.

[0025] The substantially aqueously carried product of the milling step could be added to surface coating binders dissolved or dispersed in water, solvent or mixtures of the two, to prepare a surface coating, such as an ink or paint. The reaction of certain metal flake pigments, notably aluminium, is, however unpredictable. As such a surface coating contains a proportion of water there exists the possibility that reactions may occur during storage, with the formation of hydrogen gas and associated hazards.

[0026] It is a further advantage of the process of the invention that water is substantially eliminated from contact and potential reaction with the metal pigment through the formation of the substantially non-volatile metal pigment composition.

[0027] Metals suitable for the performance of the invention include all those metals commercially employed as flake pigments, such as aluminium, zinc, copper, tin, nickel, iron and alloys thereof, such as gold bronze (an alloy of copper and zinc) or stainless steel (an alloy composed mainly of iron, nickel and chromium). Aluminium and gold bronze are preferred. There is no criticality to the particle size of the milled flakes, but a flake diameter range of 6 μm to 600 μm , preferably 10 μm to 300 μm is generally suitable. Advantageously, the particles produced are well flattened highly light reflective flake pigments.

[0028] Any compounds capable of inhibiting the reaction of the metal with water may be employed as corrosion inhibitors, for example phosphorus-, chromium-, vanadium- or silicon-containing compounds. They may

be used individually or in admixture.

[0029] Suitable phosphorus compounds may be organic or inorganic. Simple inorganic phosphates, such as calcium or magnesium phosphate, or more complex phosphosilicate compounds may be used. The latter include calcium phosphosilicate, calcium strontium phosphosilicate and aluminium zirconium zinc phosphosilicate. An especially preferred member of this class is calcium strontium zinc phosphosilicate.

[0030] Organic phosphorus compounds include alkyl and dialkyl phosphates and phosphites, with the alkyl groups containing 2-20 carbon atoms. Iso-octyl acid phosphate may be particularly mentioned. Also suitable are phosphate esters of long chain ethoxylated alcohols. Preferred amongst these is Briphos S2D, a product of the Albright & Wilson company, which is an ethoxylated phosphate ester.

[0031] Silicon containing compounds capable of inhibiting the reaction of metals with water include organic silanes and silicates, especially tetraethyl silicate, and inorganic silicon compounds such as potassium silicate.

[0032] Salts of molybdenum, vanadium and tungsten, especially the ammonium salts, have also shown particular suitability in the process of the invention.

[0033] Chromium compounds suitable for the process of the invention include ammonium dichromate and chromic acid. In this class, the former is preferred as it is less aggressive towards metal grinding media.

[0034] Amongst the above corrosion inhibiting compounds, non-resinous organic or inorganic phosphates are especially preferred. They generally offer a high level of metal passivation with few health and safety concerns.

[0035] Lubricants suitable for the process of the invention are generally organic compounds recognised in the art as having surfactant properties. Non-ionic surfactants, such as ethylene oxide condensates with aliphatic alcohols or phenols are effective without affecting the pH of the system. In general it is desirable that the pH is kept as close to neutral as possible to assist in the prevention of corrosive attack of the metal surface. Such lubricants may also be advantageous in the metal segregation and recovery stages of the process, for example to overcome the surface tension of water to ensure efficient screening. Surprisingly, fatty acids, such as oleic and stearic acids, widely used as lubricants in the solvent based ball milling of metal powders, are also effective in the milling step, especially when made active in the aqueous phase by surfactants or by formation of a water soluble alkali salt. In the same way, water insoluble plasticisers may be used, for example phthalates such as dioctyl or disodecyl phthalate, and adipates, such as dioctyl adipate.

[0036] Also suitable as lubricants are polyethylene oxides and glycols and polypropylene oxides and glycols of various molecular weights. Those displaying some water solubility are preferred.

[0037] It may also be desirable to add small quantities

of organic compounds recognised in the art for their de-foaming properties. For example, acetylenic diols may be used, but the inexpensive lower alcohols are also effective.

[0038] Typical surfactants include anionic, non-ionic and cationic surfactants, including the solid alkyl ether phosphates such as the "Crodatos CS" (Registered Trade Mark) range of Croda Chemicals Ltd, alkylaryl sulphates and their alkali metal derivatives such as alkali toluene sulphonates, alkali xylene sulphonates, alkali naphthalene sulphonates, alkali diisopropyl naphthalene sulphonates and alkali dodecyl benzene sulphonates; alcohol sulphates such as sodium lauryl alcohol sulphate, sulposuccinates such as sodium dioctyl sulposuccinate, sarcosinates such as lauroyl sarcosine and stearyl sarcosine; fatty amines such as stearylamine, and distearylamine; amine salts such as coconut fatty amine acetate; alkyl phenol ethoxylates such as nonyl phenol ethoxylate; alcohol ethoxylates such as higher ethoxylated oleyl alcohol; higher polyoxypropylene-polyoxy ethylene copolymers, such as alkylolamides such as myristic diethanolamide and coconut monoisopropanolamide, esters such as propylene glycol monostearate and cetyl palmitate; maleic anhydride copolymers such as the disodium salt of maleic anhydride and di-isobutylene, and the SMA series of low molecular weight styrene-maleic anhydride copolymers.

[0039] Phosphate esters (eg Briphos S2D) which also have lubricating and/or corrosion inhibiting properties are preferred.

[0040] Additive(s) having corrosion inhibiting, surface tension reducing and lubricating properties are present in the milling step.

[0041] Milling lubricants prevent cold welding of flakes which typically takes place during the ball milling process. The corrosion inhibitor and milling lubricant functions may be provided by a single chemical substance, for example a substituted phosphate ester (such as Briphos S2D), or the functions may be provided by two or more different materials.

[0042] There is no criticality to the mechanism of comminution. Any comminution process known in the art for metal flake production may be employed, providing the mechanical energy imparted is not so severe as to damage the metal flakes being formed. Ball milling is a widely operated process.

[0043] Neither is there any criticality to the grinding media, providing they do not react chemically with the other components during or after the comminution stage of the process. Stainless steel and high density ceramic grinding media are generally satisfactory.

[0044] Unwanted oversize or undersize metal flake particles are removed by any suitable means, such as screening of a slurry diluted by more water. The aqueous metal pigment slurry can then be concentrated to a paste by any convenient means, such as a filter press, ready for admixture with the organic binder medium.

[0045] Organic binder media include those organic

materials habitually employed as binders in paints and inks or as plastics masterbatch carriers. The organic binder chosen is dependent on the envisaged end use of the product of the invention. Thus, if desired, the binder can be a solvent-soluble resin. As examples of organic binder media there may be mentioned cellulose acetate butyrate (CAB) and cellulose acetate propionate (CAP) resins, coumarone indene, epoxy esters, epoxide melamine and epoxide phenolic condensates, ketone, aldehyde, maleic and phenolic resins and also rosin, cellulose and petroleum derived resins, together with thermoplastic polymers, such as polyacrylates, polyolefins, polyvinyls, styrenics, polyamides, polyesters and copolymers thereof. Also suitable are natural and synthetic waxes, such as montan and paraffin wax and synthetic waxes such as polyethylene and polypropylene waxes. Where it is desired to introduce liquid organic binders, such as plasticisers, for example to improve application properties, they are desirably added in minor proportions in combination with solid resins, to provide structural rigidity in the resulting metal pigment particles. Such plasticisers include mineral oils as well as phthalates such as dioctyl or diisodecyl phthalate, and adipates such as dioctyl adipate.

[0046] Gelatin and carrageenan are useful water soluble organic binders due to their suitability for food contact applications.

[0047] In a preferred embodiment the binder material is a non-resinous material.

[0048] The products of the invention are typically in a form, such as tablet, pellet, granule, flake or spherical bead, which provides the attributes of ease of handling, low- or non-dusting and meterability. Granules having a length of 5-20mm, a cylindrical cross section and a diameter of 1.5-3mm are preferred, as they have been found to offer optimum handling characteristics, especially in plastics applications where it is important to prevent stratification of polymer pellets and the particularly dense products of the invention. Apparatus used for the preparation of such physical forms is well known to those skilled in the art and is described in, for example, European Patent 0134676.

[0049] The function of the organic binder medium is to bind the metal pigment flakes together to prevent them becoming airborne as dust. Where the organic binder medium is soluble or dispersible in water, it may be added as an aqueous solution or dispersion. Where it is insoluble in water it is preferable to micronise the binder. In this latter case, the coherent paste of metal pigment, water and micronised organic binder medium is formed into particles, for example by mixing and extrusion, then dried at for example 60°C to remove substantially all the water, then heat treated at elevated temperature, for example 100 to 130°C for a short time, to fuse together the micronised organic binder medium and the metal pigment. This process is particularly suited to products for the plastics market, in which suitable organic binder media tend to be insoluble in water. It

provides a final product form which has excellent colouristic and application properties when used in the injection moulding or extrusion of thermoplastics.

[0050] In extreme cases the chosen binder may be insoluble in water and unsuitable for micronising and fusion. In such cases the binder may be dissolved or dispersed in a suitable organic solvent prior to admixture with the metal pigment component. Such a solvent is desirably, though not essentially, water miscible.

[0051] In a preferred embodiment the metal is aluminium. Conveniently the lubricant is a mixture of a surfactant and a fatty acid, such as oleic or stearic acid. Oleic acid in an amount of 1 to 100%, preferably 2 to 60%, most preferably 3 to 8%, for example approximately 5%, by weight relative to the metal has been found to be effective in mixture with non-ionic surfactants at 5-15%. The corrosion inhibitor may advantageously be a phosphate ester having a polyethylene oxide side chain.

[0052] The invention is further illustrated by the following Examples in which all parts and percentages are by weight, unless otherwise stated.

Example 1

[0053] To a cylindrical mill of diameter 21 cm were added:

5000g of 1/8 inch (3.2mm) diameter stainless steel balls;
500g of distilled water;
100g of fine particle size, aluminium powder; and
13g of Briphos S2D, which had been pre-neutralised with triethanolamine - this makes the Briphos more soluble and reduces attack of the aluminium. Briphos S2D is a phosphated ester of a long chain ethoxylated alcohol manufactured by Albright and Wilson. The mill was rotated at 80 rpm for 4 hours and the aluminium pigment so formed separated from the steel balls by sieving, prior to recovery in a filter funnel.

[0054] In a laboratory mixer, a quantity of the aluminium flake pigment thus obtained, containing 25.0g solids, was added to 10.7g Hoechst Ceridust 3620, a micronised polyolefin wax with a particle size of less than 22 microns. The mixture was formed into granules by extrusion through a 2mm diameter die, and dried in an oven held at a temperature of 60°C +/- 10°C for two hours. The granules were then placed in an oven held at 115°C +/- 10°C for 10 minutes to fuse the organic binder medium. 35.1g of substantially water free, non-dusting, free flowing, meterable granules of approx. 12mm length and of good abrasion resistance were obtained.

[0055] A test piece made by injection moulding the granule in polystyrene at a level of 1% calculated on polymer weight, had a smooth, fine and bright silvery ap-

pearance.

Example 2

5 [0056] The following mixture was milled in a cylindrical mill as described in Example 1, with the following formulation:-

10 5000g of 1/8 inch (3.2mm) diameter stainless steel balls;
500g of distilled water;
100g of aluminium powder of approx. 33µm average particle diameter;
13g of Briphos S2D, which had been neutralized
15 with triethanolamine;
2.85g of Antarox V27, a linear C₈₋₁₀ aliphatic alcohol; ethoxylate/propoxylate alcohol manufactured by Rhone-Poulenc Chemicals; and
0.15g of Surfynol 104, an acetylinic diol from Air
20 Products.

[0057] A quantity of the derived aluminium flake pigment paste, containing 24.1g solids, was added to 10.3g of FE 532-00, a micronised ethylene vinyl acetate resin
25 sold by Croxton and Garry. The mixture was granulated, dried and heat treated as in Example 1. A test piece made by injection moulding the granule in polypropylene at a level of 3%, calculated on polymer weight, had a bright, highly sparking, silvery appearance.

Example 3

[0058] The method of Example 1 was repeated with the substitution of Hoechst Ceridust 3620 by an equal
35 weight of Polyox WSR 301, a solid, non-ionic, high molecular weight, water-soluble, polyethylene oxide polymer. The mixture was formed into granules by extrusion through a 2mm diameter die, and dried in an oven held at a temperature of 50°C +/- 10°C for four hours. 35.2g
40 of substantially water free, non-dusting, free flowing, meterable granules of approx. 12mm length and of good abrasion resistance were obtained.

[0059] An aqueous printing ink was prepared by combining the following:-

45 20g of the granule product of this Example, dispersed in 25g water and let down with
53g Zinpol 132, an acrylic resin emulsion of 40% solids, supplied by Worlee Chemie and
50 2g waxes and defoaming additives.

The ink thus formed was stable to gassing and provided a smooth, bright metallic silver effect when printed on a paper substrate.

Example 4

[0060] 10g Halox ZSP 391, a calcium strontium zinc

phosphosilicate compound, sold by Halox Pigments of Hammond, Indiana, United States of America.

3g oleic acid,
5g Pluriol 600, a polypropylene glycol supplied by BASF A.G., and
50g deionised water

were high speed stirred to solution.

35.1g of solution,
219.4g deionised water, and
30g atomised aluminium powder of approx. 8µm average particle diameter

were set on a vibratory mill with 1500g 1/8 inch (3.2mm) diameter stainless steel balls, in a rigid plastic pot. After 4 hours, the contents were removed and separated from the steel balls. Water was added to a metal concentration of about 3% and the slurry screened on a 400 mesh screen. The undersize fraction was concentrated in a filter press to about 80% metal content, then combined with Microthene 532-00, an ethylene vinyl acetate polymer sold by Croxton & Garry Ltd, by the fusion process of Example 1.

[0061] When injection moulded at 1.5% w/w in polypropylene, the granule exhibited a particularly bright, reflective silver coloration, with good dispersion.

[0062] The dispersibility characteristics of this product can be further enhanced by replacing up to 30% of the Microthene by a high boiling mineral oil such as Kaydol, manufactured by Witco.

Example 5

[0063]

200g Halox SZP-391
100g Pluriol P600
100g Antarox V27
100g water and
60g oleic acid

were high speed stirred for 10 minutes to form a homogeneous mixture. To a cylindrical mill of diameter 21cm were added: 5000g of 1/8 inch (3.2mm) diameter stainless steel balls, 75g of the above mixture, 100g aluminium powder of 9µm median particle diameter and 375g of distilled water.

[0064] The mill was rotated at 105 rpm for 6 hrs to form a flake pigment product of approximately 25µm median particle diameter. The grinding media were removed on a coarse screen and the product itself screened on a 400 mesh (38µm) screen using water as the washing liquid. The flake passing through the screen was concentrated on a filter funnel. When converted to a granule and tested in the water based ink system of Example 3, this material provided a bright silver effect

with good coverage and good adhesion to a paper substrate.

[0065] The Halox SZP-391 of this Example may be replaced by members of the Heucophos range of multiple metal phosphates manufactured by Heubach GmbH, with similar results.

Example 6

[0066]

19g Antarox V27 and
1g Surfynol 104

were warmed and mixed together to form a homogeneous liquid, which was mixed into 120g of the water based paste intermediate of Example 1, containing 80g metal, to form a stiff paste. The paste was extruded through a 2.5mm diameter die into strands and the water removed by drying at 70°C. The resulting dry strands were broken up into granules of approximately 1cm length. An aqueous ink was prepared by predispersing the granules in a small quantity of water into which the Zinpol medium of Example 3 was thereafter blended. A bright ink, with good stability and coverage was obtained.

[0067] The Antarox and Surfynol in this example may be replaced by 20g of the natural product carageenan in powder form with similar results. Such a granule has particularly good shelf life and redispersion in water.

Example 7

[0068] To a 0.5 litre capacity vibratory pot mill were added:-

1.5kg of 2.4mm diameter ceramic grinding media,
150g water,
30g aluminium powder of 7µm average particle diameter,
2.5g Iso-octyl acid phosphate and sufficient 25% aqueous ammonia to maintain the pH at approximately 7.

[0069] The charge was milled for 3 hrs then recovered, screened, formed into granules and tested as in Example 5. The printed ink had similar brightness, but superior coverage to that of Example 5.

Example 8

[0070] To 0.5 litre capacity vibratory pot mill were added:-

1.5kg of 3mm diameter stainless steel grinding media, 150g water,
30g aluminium powder of 7µm average particle diameter,
2.4g Briphos S2D and

0.6g 25% aqueous ammonia.

[0071] The charge was milled for 4 hrs, then discharged with water and screened on a 400 mesh (38 μ m) screen. The fraction passing through the screen was collected and concentrated under vacuum in a filter funnel. A quantity of the filter cake, corresponding to 80g metal was thoroughly mixed with 20g of a commercial source of C₁₂₋₁₄-aliphatic alcohols. The stiff paste was formed into pellets and the water removed at 65°C. The resulting product had good stability and colouristic properties in a one pack UV curing resin system.

Example 9

[0072] The milled, screened and recovered paste product of Example 8 was mixed with micronised Laropal K80 ketone resin, manufactured by BASF AG in the ratio 80 parts by weight of metal to 20 parts by weight of resin. The mixture was formed into granules by extrusion and the water removed in an oven of 70°C.

[0073] When tested in a solvent based ink, the granular product displayed similar characteristics to a granule prepared by the methods described in EP-A-0134676 using a conventional solvent milled paste as starting material.

Example 10

[0074] An amount of the milled, screen and recovered paste product of Example 8 corresponding to 80g metal was placed in a 1 litre beaker fitted with a propeller agitator set at 400 rpm. With agitation there were then added,

530g Dowanol PB29, an ether-alcohol solvent of the Dow Chemical Co,
26.7g of 25 % aqueous ammonia,
20g tetraethyl silicate and finally another
26.7g of 25% aqueous ammonia.

The mixture was agitated for 5 minutes, then there was added another 26.7g of 25% aqueous ammonia.

[0075] After a further two hours mixing, air was bubbled through the slurry for 2 hours to remove excess ammonia and the metal recovered on a Buchner filter. Thereafter, the filter cake was converted to a gold pigment by the process of Example 1 of our co-pending British Patent Application No 9809507.0 (ie was stirred at 600 rpm with a paddle agitator for one hour, then filtered under vacuum on a Buchner funnel, and washed with excess water. The product was a viscous paste dispersible in both water and oxygen-containing hydrocarbon solvents) and then to a granule by the process of Example 25 of EP-A-0134676. The granules so formed provided a rich gold coloration in water based paints and inks.

Example 11

[0076]

2g of oleic and
2g of Antarox V27 and Surfynol 104 in 19:1 weight ratio were mixed together and added dropwise to 140g vigorously stirred water in a beaker containing 1g isopropanol, to form an emulsion. There was then added
4g hydrogen peroxide (30%w/w; 100 volume) and the whole added to
30g A 7401 atomised aluminium powder of 8 μ m average particle diameter in a ball milling pot of 500ml volume containing
1500g spherical, stainless steel grinding media of 3mm average diameter.

The pot was sealed and placed on a vibratory mill for 4 hours. The charge was washed out with 200g water, screened on a 400 mesh screen and an aqueous paste recovered by vacuum filtration.

The water was removed by drying at 50°C and 16g of the resulting fine flake powder incorporated into 8.89g Worleesol 31, a modified linseed oil polymer (supplied by Worlee Chemie as a 45% solution in 80/20 w/w water/2-butoxy ethanol), further diluted in 3.5g of a 80/40 weight ratio mixture of water and isopropanol. After forming into granules and evaporating the volatiles at 60°C, the soft, dry product was found to give a bright, high coverage silver finish in an exclusively isopropanol based or water/isopropanol based varnish.

Example 12

[0077] A formulation comprising

1500 spherical, ceramic grinding media of 3mm average diameter,
150g water,
30g aluminium powder of 8 μ m average particle diameter,
2.4g Crodafos 25D5A, an ethoxy (5) C₁₂₋₁₅ alcohol acid phosphate ester, supplied by Croda Ltd and
0.8g ammonia, 25% in water was loaded into a 500ml pot and milled on a vibratory mill for 3 hours.

The grinding media was removed on a coarse screen and the metal flake pigment product screened on a 400 mesh (38 μ m) screen using water as the washing liquid. The flake passing through the screen was concentrated on a filter funnel. When converted to a granule with a carrageenan carrier and tested in the water based ink system of Example 3, this material provided a bright silver effect with good coverage.

Example 13

[0078] The method of Example 12 was repeated with the substitution of Crodafos 25D5A by an equal weight of Crodafos T5A, an ethoxy (5) isotridecanol acid phosphate ester. Similar results were obtained on conversion to a granule and testing as described therein.

Example 14

[0079] The method of Example 12 was repeated with the substitution of Crodafos 25D5A by an equal weight of Crodafos N10N, an ethoxy (10) oleyl alcohol phosphate ester diethylamine salt. Similar colouristic results were obtained on conversion to a granule and testing as described therein. This material, however, also had outstanding adhesion to the paper substrate, as demonstrated by a tape test.

Example 15

[0080] A formulation comprising

1500 spherical, stainless steel grinding media of 3mm average diameter,
150g distilled water,
30g aluminium powder of 25µm average particle diameter and
2g octyl phosphonic acid

was charged to a 500ml pot and milled on a vibratory mill for 1.5 hours.

After removing the grinding media, the milled material was screened on a 325 mesh (45µm) screen and recovered as a granule by the method of Example 12 using as carrier a water soluble part hydrolysed polyvinyl acetate resin. The 80% metal content granule provided a very bright, sparking coating based on Joncryl 537, an aqueous acrylic industrial paint resin manufactured by S C Johnson.

Example 16

[0081] Aluminium pigment, made by the process of example 1 as a water carried paste containing 40g of aluminium metal, was mixed with 13.3 g of Beette BE 370, hexamethoxymethyl melamine from BIP Speciality Resins Ltd. The mixture was formed into granules by extrusion through a 2 mm diameter die, and dried in an oven held at a temperature of 60°C for 16 hours.

[0082] An aqueous printing ink was made by dispersing 5 g of the dried granule in 5 g of distilled water, and then mixing this dispersion with 15 g of Glascol LS2, a colourless water based ink from Allied Colloids. A draw-down on to white paper showed that the product had a bright metallic silver colour.

Example 17

[0083] The procedure of example 16 was repeated with the Beette BE 370 replaced by RC-294-J427, an acetylenic diol surfactant from Air Products and Chemicals Inc. The dried granular product so formed gave a brilliant metallic silver effect when made up into a Glascol LS2 ink.

Example 18

[0084] A dispersion of

7.5 g Molywhite 212, a basic calcium zinc molybdate from Sherwin Williams Chemicals
1.5g oleic acid
2.5g Pluriol P600
2.5 g Antarox V27
25.0 g distilled water

was added to

30 g of a fine particle size aluminium powder
100 g distilled water
1500 g of 1/8 inch diameter stainless steel balls

and milled on a vibratory mill for 3.5 hours.

[0085] A flaked aluminium paste product was extracted from the grinding media. If required, the paste can be converted into granular form as described in previous Examples.

Claims

1. A process for preparing a low- or non-dusting, substantially non-volatile metal flake pigment composition, said process comprising ball milling atomised metal powder in the presence of a milling fluid, together with one or more additive(s) which together exhibit lubricating, corrosion inhibiting and surfactant properties, the lubricant being a non-suet lubricant; wherein said milling fluid consists substantially of water and has a maximum content of organic solvent of 10% by weight.
2. A process as claimed in claim 1 which includes the step of screening the milled composition to select particles of the desired size.
3. A process as claimed in either one of claims 1 and 2 which includes the step of forming a coherent paste of an organic binder medium, water and metal flake pigment.
4. A process for preparing a low- or non-dusting, substantially non-volatile metal flake pigment composition, said process comprising:

- a) ball milling atomised metal powder in the presence of a milling fluid, together with either an additive having lubricating, corrosion inhibiting and surfactant properties or with a lubricant and at least one corrosion inhibitor, wherein said milling fluid consists substantially of water; and
- b) forming a coherent past of an organic binder medium, water and metal flake pigment, wherein said paste is formed by mixing a first component comprising organic binder medium and a second component comprising metal pigment, with either or both of the first and second components comprising water and the paste containing from 3-70% of the organic binder medium based on the weight of the metal pigment, and either sub-dividing the coherent paste into particles and removing substantially all volatile liquids from the particles, or removing substantially all volatile liquids from the coherent paste and sub-dividing the resulting mass into particles, at least 98% by weight of the resulting particles being retained on a sieve having a 150µm aperture and each containing a plurality of metal pigment particles dispersed in a matrix of organic binder medium, the organic binder medium being capable of binding the metal flake pigment particles by either precipitation from solution during volatile liquid removal, or sintering or melting at elevated temperature and fusion.
5. A process as claimed in any one of claims 1 to 4 wherein said lubricant comprises an ethylene oxide condensate with alcohols or phenols, and a fatty acid or a derivative thereof.
 6. A process as claimed in any one of claims 1 to 5 wherein said corrosion inhibitor is a phosphorus-, chromium-, vanadium- or silicon-containing compound, or calcium zinc molybdate.
 7. A process as claimed in claim 6 wherein said corrosion inhibitor is a phosphate ester having a polyethylene oxide side chain.
 8. A process as claimed in any one of claims 1 to 4 wherein a single moiety acts as lubricant and corrosion inhibitor.
 9. A process as claimed in claim 8 wherein said moiety is a phosphate ester having a polyethylene oxide side chain.
 10. A process as claimed in any one of claims 1 to 9 wherein a surfactant is present in said milling step.
 11. A process as claimed in any one of claims 1 to 10 wherein the metal pigment is of aluminium, zinc, copper, tin, nickel, iron or alloys thereof.
 12. A process as claimed in any one of claims 1 to 11 wherein the metal pigment is of aluminium or gold bronze.
 13. A process as claimed in any one of Claims 1 to 12 wherein the metal pigment is a highly light reflective pigment.
 14. A process as claimed in any one of claims 1 to 13 wherein the metal pigment is aluminium, the lubricant is oleic acid and the corrosion inhibitor is a phosphate ester having a polyethylene oxide side chain.
 15. A process as claimed in any one of claims 1 to 14 which includes a step of treating the milled metal pigment flakes in an aqueous system.
 16. A process as claimed in claim 15 wherein said milled metal pigment flakes are treated with chromate, coated with silica or coated with alumina.
 17. A process as claimed in any one of claims 1 to 15 wherein said pigments are coated with micronised binder and sintered.
 18. Use of a phosphate ester having a polyethylene oxide side chain as an additive in a process of ball milling a metal powder as claimed in any one of Claims 1 to 17.
 19. A paste comprising a ball-milled metal flake pigment, water and agent(s) having lubricating and corrosion inhibiting properties, said paste being obtainable from a process as claimed in any one of Claims 1 to 17.
 20. A paste as claimed in claim 19 comprising ball-milled aluminium or gold bronze flake pigment, water and a phosphate ester having a polyethylene oxide side chain.
 21. A granule containing a plurality of milled metal pigment particles obtainable from a process as claimed in any one of Claims 1 to 17, dispersed in a matrix of organic binder medium which is substantially free of organic solvent.
 22. A granule as claimed in claim 21 which is a sintered granule and wherein said metal pigment particles are dispersed in a matrix of micronised organic binder medium.

Patentansprüche

1. Ein Verfahren zur Herstellung einer gering- oder nichtstaubenden, im wesentlichen nichtflüchtigen Metallflockenpigmentzusammensetzung, wobei das Verfahren das Mahlen von zerstäubtem Metallpulver mittels einer Kugelmühle unter Vorhandensein einer Mahflüssigkeit umfaßt, zusammen mit einem oder mehreren Zusatzstoffen, die zusammen schmierende, korrosionshemmende und grenzflächenaktive Eigenschaften aufweisen, wobei das Schmiermittel ein nichttaliges Schmiermittel ist; wobei die Mahflüssigkeit im wesentlichen aus Wasser besteht und einen maximalen Gehalt an organischen Lösungsmitteln von 10 Gew.-% aufweist. 5
2. Verfahren gemäß Anspruch 1, welches den Schritt des Siebens der gemahlten Zusammensetzung, um Partikel der gewünschten Größe auszuwählen, umfaßt. 10
3. Verfahren gemäß Anspruch 1 und 2, welches den Schritt des Bildens einer kohärenten Paste aus einem organischen Bindemittel, Wasser und Metallflockenpigmenten umfaßt. 15
4. Verfahren zur Herstellung einer gering- oder nichtstaubenden nichtflüchtigen Metallflockenpigmentzusammensetzung, wobei das Verfahren folgendes umfaßt: 20
 - a) Mahlen von zerstäubtem Metallpulver mittels einer Kugelmühle unter Vorhandensein einer Mahflüssigkeit, zusammen mit entweder einem Zusatzstoff, der schmierende, korrosionshemmende und grenzflächenaktive Eigenschaften aufweist, oder mit einem Schmiermittel und mindestens einem Korrosionshemmstoff, wobei die Mahflüssigkeit im wesentlichen aus Wasser besteht; und 25
 - b) Bilden einer kohärenten Paste aus einem organischen Bindemittel, Wasser und Metallflockenpigmenten, wobei die Paste durch Mischen einer ersten Komponente, die organisches Bindemittel umfaßt, und einer zweiten Komponente, die Metallpigment umfaßt, wobei entweder eine oder beide der ersten und zweiten Komponenten Wasser enthalten und wobei die Paste 3-70 % des organischen, auf dem Gewicht des Metallpigments basierenden Bindemittels enthält, und entweder Unterteilen der kohärenten Paste in Partikel und Entfernen von im wesentlichen allen flüchtigen Flüssigkeiten von den Partikeln, oder Entfernen von im wesentlichen allen flüchtigen Flüssigkeiten von der kohärenten Paste und Unterteilen der verbleiben- 30
- den Masse in Partikel, wobei mindestens 98 Gew.-% der resultierenden Partikel in einem Sieb mit der Öffnung 150 µm zurückbleiben und jedes eine Vielzahl von Metallpigmentpartikeln, die in einer Matrix aus organischem Bindemittel gelöst sind, enthält, wobei das organische Bindemittel die Metallflockenpigmentpartikel durch entweder Ausfällen aus der Lösung während dem Entfernen der flüchtigen Flüssigkeiten oder durch Sintern oder Schmelzen bei erhöhter Temperatur und Fusion binden kann. 35
5. Verfahren gemäß einem der Ansprüche 1 bis 4, wobei das Schmiermittel ein Ethylenoxidkondensat mit Alkoholen oder Phenolen und eine Fettsäure oder ein Derivat davon umfaßt. 40
6. Verfahren gemäß einem der Ansprüche 1 bis 5, wobei der Korrosionshemmstoff eine Phosphor, Chrom, Vanadium oder Silizium enthaltende Mischung ist. 45
7. Verfahren gemäß Anspruch 6, wobei der Korrosionshemmstoff ein Phosphatester mit einer Polyethylenoxid-Seitenkette ist. 50
8. Verfahren gemäß einem der Ansprüche 1 bis 4, wobei eine einzelne Komponente als Schmiermittel und Korrosionshemmstoff wirkt. 55
9. Verfahren gemäß Anspruch 8, wobei die Komponente ein Phosphatester mit einer Polyethylenoxid-Seitenkette ist.
10. Verfahren gemäß einem der Ansprüche 1 bis 9, wobei ein grenzflächenaktiver Stoff in dem Mahschritt vorhanden ist.
11. Verfahren gemäß einem der Ansprüche 1 bis 10, wobei das Metallpigment aus Aluminium, Zink, Kupfer, Zinn, Nickel, Eisen oder Legierungen davon besteht.
12. Verfahren gemäß einem der Ansprüche 1 bis 11, wobei das Metallpigment aus Aluminium oder Goldbronze besteht.
13. Verfahren gemäß einem der Ansprüche 1 bis 12, wobei das Metallpigment ein stark lichtreflektierendes Pigment ist.
14. Verfahren gemäß einem der Ansprüche 1 bis 13, wobei das Metallpigment Aluminium, das Schmiermittel Oleinsäure und der Korrosionshemmstoff ein Phosphatester mit einer Polyethylenoxid-Seitenkette ist.
15. Verfahren gemäß einem der Ansprüche 1 bis 14,

weiches einen Schritt des Behandelns der gemahlenden Metallpigmentflocken in einem Wassersystem beinhaltet.

16. Verfahren gemäß Anspruch 15, wobei die gemahlenden Metallpigmentflocken mit Chrom behandelt werden und mit Silika oder mit Aluminiumoxid beschichtet werden.
17. Verfahren gemäß einem der Ansprüche 1 bis 15, wobei die Pigmente mit mikronisiertem Bindemittel beschichtet und gesintert werden.
18. Verwenden eines Phosphatesters mit einer Polyethylenoxid-Seitenkette als Zusatzstoff in einem Vorgang des Mahlens eines Metallpulvers mittels einer Kugelmühle wie in jedem der Ansprüche 1 bis 17 beansprucht.
19. Eine Paste, die ein mittels einer Kugelmühle gemahlendes Metallflockenpigment, Wasser und Stoff (e) mit schmierenden und korrosionshemmenden Eigenschaften umfaßt, wobei die Paste durch ein Verfahren gemäß den Ansprüchen 1 bis 17 gewonnen wird.
20. Paste gemäß Anspruch 19, bestehend aus einem mittels einer Kugelmühle gemahlenden Aluminium oder Goldbronzeflockenpigment, Wasser und einem Phosphatester mit einer Polyethylenoxid-Seitenkette.
21. Eine Granalie, bestehend aus einer Vielzahl an gemahlenden Metallpigmentpartikeln, die durch ein Verfahren gemäß der Ansprüche 1 bis 17 gewonnen werden, aufgelöst in einer Matrix eines organischen Bindemittels, die im wesentlichen frei von organischem Lösungsmittel ist.
22. Granalie gemäß Anspruch 21, welche eine gesinterte Granalie ist und wobei die Metallpigmentpartikel in einer Matrix von mikronisiertem organischen Bindemittel gelöst sind.

Revendications

1. Un procédé de préparation d'une composition à pigment métallique en écailles sensiblement non volatile et formant peu ou pas de poussière, ledit procédé comprenant le broyage par billes de poudre métallique atomisée en présence d'un fluide de broyage ainsi que d'un ou plusieurs additifs, lesquels exhibent ensemble des propriétés lubrifiantes, inhibitrices de corrosion et tensioactives, le lubrifiant n'étant pas un lubrifiant à base de graisse de boeuf ; dans lequel ledit fluide de broyage est sensiblement composé d'eau et possède une te-

neur maximale en solvant organique de 10 % en poids.

2. Un procédé tel que revendiqué dans la revendication 1, lequel comporte l'étape consistant à cribler la composition broyée pour sélectionner des particules de la taille désirée.
3. Un procédé tel que revendiqué dans l'une ou l'autre des revendications 1 et 2, lequel comporte l'étape consistant à former une pâte cohérente avec un milieu liant organique, de l'eau et du pigment métallique en écailles.
4. Un procédé de préparation d'une composition à pigment métallique en écailles sensiblement non volatile et formant peu ou pas de poussière, ledit procédé comprenant :
 - (a) le broyage par billes de poudre métallique atomisée en présence d'un fluide de broyage ainsi que soit d'un additif possédant des propriétés lubrifiantes, inhibitrices de corrosion et tensioactives, soit d'un lubrifiant et au moins un inhibiteur de corrosion, dans lequel ledit fluide de broyage est sensiblement composé d'eau ; et
 - (b) la formation d'une pâte cohérente avec un milieu liant organique, de l'eau et du pigment métallique en écailles, dans lequel ladite pâte est formée en mélangeant un premier composant comprenant du milieu liant organique et un second composant comprenant du pigment métallique, le premier ou le second composant, ou les deux, comprenant de l'eau et la pâte contenant de 3 à 70 % du milieu liant organique rapporté au poids du pigment métallique, et soit en subdivisant la pâte cohérente en particules et en éliminant sensiblement tous les liquides volatils des particules, soit en éliminant sensiblement tous les liquides volatils de la pâte cohérente et en subdivisant la masse résultante en particules, 98 % en poids au moins des particules résultantes étant retenues sur un tamis ayant une ouverture de 150 µm et chacune contenant une pluralité de particules de pigment métallique dispersées dans une matrice de milieu liant organique, le milieu liant organique étant à même de lier les particules de pigment métallique en écailles soit par précipitation à partir de la solution pendant l'élimination des liquides volatils, soit par frittage, soit en les faisant fondre à une température élevée et par fusion.
5. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 4 dans lequel ledit lu-

brillant comprend un produit de condensation d'oxyde d'éthylène avec des alcools ou des phénols et un acide gras ou un dérivé de celui-ci.

6. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 5 dans lequel ledit inhibiteur de corrosion est un composé contenant du phosphore, du chrome, du vanadium ou du silicium. 5
7. Un procédé tel que revendiqué dans la revendication 6 dans lequel ledit inhibiteur de corrosion est un ester phosphate possédant une chaîne latérale d'oxyde de polyéthylène. 10
8. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 4 dans lequel un fragment unique fait office de lubrifiant et d'inhibiteur de corrosion. 15
9. Un procédé tel que revendiqué dans la revendication 8 dans lequel ledit fragment est un ester phosphate possédant une chaîne latérale d'oxyde de polyéthylène. 20
10. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 9 dans lequel un tensioactif est présent dans ladite étape de broyage. 25
11. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 10 dans lequel le pigment métallique est en aluminium, zinc, cuivre, étain, nickel, fer ou en alliages de ceux-ci. 30
12. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 11 dans lequel le pigment métallique est en aluminium ou en bronze doré. 35
13. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 12 dans lequel le pigment métallique est un pigment hautement réflecteur de lumière. 40
14. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 13 dans lequel le pigment métallique est de l'aluminium, le lubrifiant est de l'acide oléique et l'inhibiteur de corrosion est un ester phosphate possédant une chaîne latérale d'oxyde de polyéthylène. 45
15. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 14, lequel comporte une étape consistant à traiter les écailles de pigment métallique broyé dans un système aqueux. 50
16. Un procédé tel que revendiqué dans la revendication 15 dans lequel lesdites écailles de pigment métallique broyé sont traitées avec du chromate, en-

duites de silice ou enduites d'alumine.

17. Un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 15 dans lequel lesdits pigments sont enduits de liant micronisé et frittés. 5
18. Utilisation d'un ester phosphate possédant une chaîne latérale d'oxyde de polyéthylène en tant qu'additif dans un procédé de broyage par billes d'une poudre métallique tel que revendiqué dans n'importe laquelle des revendications 1 à 17. 10
19. Une pâte comprenant un pigment métallique en écailles broyé par billes, de l'eau et un ou des agents possédant des propriétés lubrifiantes et inhibitrices de corrosion, ladite pâte pouvant être obtenue à partir d'un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 17. 15
20. Une pâte telle que revendiquée dans la revendication 19 comprenant du pigment en écailles d'aluminium ou de bronze doré broyé par billes, de l'eau et un ester phosphate possédant une chaîne latérale d'oxyde de polyéthylène. 20
21. Un granulé contenant une pluralité de particules de pigment métallique broyé pouvant être obtenues à partir d'un procédé tel que revendiqué dans n'importe laquelle des revendications 1 à 17, dispersées dans une matrice de milieu liant organique, laquelle est sensiblement exempte de solvant organique. 25
22. Un granulé tel que revendiqué dans la revendication 21, lequel est un granulé fritté et dans lequel lesdites particules de pigment métallique sont dispersées dans une matrice de milieu liant organique micronisé. 30